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1: Wiad Lek 1990 Jan 1-15;43(1-2):4-9

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[Angiogenic activity of mononuclear cells in the peripheral blood of patients with ischemic heart disease]

[Article in Polish]

Pawinska M, Laniewska I, Sztorc M, Syska J, Wroblewski T, Kuch J.

Katedry i Kliniki Kardiologii II Wydziału Lekarskiego Ak. Med., Warszawie.

In the paper the cell-mediated phase of the immune response was assessed in ischaemic heart disease (IHD). The assessment was based on the angiogenesis test in which new capillaries are formed from the already existing vascularization. The process is mediated by mononuclear cells from peripheral blood, and is induced by degradation products appearing as a result of ischaemic injury to the tissues. The test was carried out in 71 patients with IHD and in 65 clinically healthy subjects. A significant fall was demonstrated of the angiogenic activity of mononuclear cells isolated from the peripheral blood of IHD patients as compared with controls (p less than 0.001). No effect was method of disease duration, its form or treatment on angiogenesis. The obtained results may suggest a failure of the immune system competent in this process in IHD. Perhaps this is related to a special form of IHD.

PMID: 1695042 [PubMed - indexed for MEDLINE]

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NEWS 44 Feb 24 PCTGEN now available on STN

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NEWS 51 Mar 24 PATDPAFULL now available on STN

NEWS 52 Mar 24 Additional information for trade-named substances without structures available in REGISTRY

NEWS 53 Mar 24 Indexing from 1957 to 1966 added to records in CA/CAPLUS

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*> s bone (a) marrow

L1 617302 BONE (A) MARROW

*> s 11 and mononuclear (a) cell

2 FILES SEARCHED...

6 FILES SEARCHED...

L2 23479 L1 AND MONONUCLEAR (A) CELL

*> s 12 and angiogenesis?

L3 1465 L2 AND ANGIOGENES?

SEARCH ENDED BY USER

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*> s 12 and vessel?

L4 2771 L2 AND VESSEL?

*> dup rem 13

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L5 1393 DUP REM L3 (72 DUPLICATES REMOVED)

*> s 15 not PY=>2000

2000 NOT A VALID FIELD CODE

5 FILES SEARCHED...

L6 211 L5 NOT PY=>2000

*> d 1-20

L6 ANSWER 1 OF 211 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

AN 2000:24:767 BIOSIS

DN PREV200000024:767

TI Local transplantation of autologous bone marrow

-derived mononuclear cells augments collateral vessel

formation in ischemic hindlimb in rabbits.

AU Shintani, Sachiko (1); Murohara, Toyoki; Ueno, Takafumi; Ikeda, Hisao;

Duan, Junji; Imazumi, Tetsuro (1) Kurume Univ., Kurume, Japan

CS Circulation, (Nov. 2, 1999) Vol. 110, No. 18 SUPPL., pp. 1-406.

SO Meeting Info.: 72nd Scientific Sessions of the American Heart Association

Atlanta, Georgia, USA November 7-10, 1999

ISSN: 0009-7322.

DT Conference

LA English

FS German

LA German

CS HAYASHIBARA BIOCHEM LABS INC., FUJISAKI CELL CTR, OKAYAMA, JAPAN (Reprint); HAYASHIBARA BIOCHEM LABS INC., FUJISAKI INST., OKAYAMA, JAPAN

CVA JAPAN CELLULAR IMMUNOLOGY, (15 DEC 1999) VOL. 198, NO. 2, PP. 111-119. Publisher: ACADEMIC PRESS INC, 525 B ST, STE 1900, SAN DIEGO, CA 92101-4495. ISSN: 0008-8739.

DT Article; Journal

FS LIFE

LA English

REC Reference Count: 27

*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS.

*ANSWER 4 OF 211 SCISEARCH COPYRIGHT 2003 ISI (R)

LA 1991:385381 SCISEARCH

AN The Genuine Article (R) Number: 195KH

TI Blast cell-surface and plasma soluble urokinase receptor in acute leukemia patients: Relationship to classification and response to therapy

AU Mustjoki S (Reprint); Alitalo K; Stephens R W; Vaheki A

CS UNIV HELSINKI, HAARTMAN INST., DEPT VIROL, POB 21, FIN-00014 HELSINKI, FINLAND (Reprint); UNIV HELSINKI, HAARTMAN INST, CENT HOSP, DEPT MED, DIV HAEMATOL, HELSINKI, FINLAND; RIGSHOSP, FINSEN INST, DK-2100 COPENHAGEN, DENMARK

CVA FINLAND; DENMARK

SO THROMBOSIS AND HAEMOSTASIS, (MAY 1999) VOL. 81, NO. 5, PP. 705-710.

Publisher: F. SCHATTNER VERLAG GMBH, P O BOX 10 45 45, LENZHALDE 3,

D-70040 STUTTGART, GERMANY.

ISSN: 0340-6245.

DT Article; Journal

FS LIFE

LA English

REC Reference Count: 37

*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS.

L6 ANSWER 5 OF 211 SCISEARCH COPYRIGHT 2003 ISI (R)

LA 1991:284019 SCISEARCH

AN The Genuine Article (R) Number: 183VC

TI Ischemia- and cytokine-induced mobilization of bone

marrow-derived endothelial progenitor cells for neovascularization

AU Takahashi T; Kaku C; Masuda H; Chen D; Silver M; Kearney M; Mgner M; Tener J M (Reprint); Asahara T

CS TUFTS UNIV, ST ELIZABETHS MED CTR, SCH MED, DEPT MED CARDIOL, 736 CAMBRIDGE ST, BOSTON, MA 02135 (Reprint); TUFTS UNIV, ST ELIZABETHS MED CTR, SCH MED, DEPT MED CARDIOL, BOSTON, MA 02135; TUFTS UNIV, ST ELIZABETHS MED CTR, SCH MED, DEPT BIOMED RES, BOSTON, MA 02135

CVA USA

SO NATURE MEDICINE, (APR 1999) VOL. 5, NO. 4, PP. 434-438.

Publisher: NATURE AMERICA INC, 345 PARK AVE SOUTH, NEW YORK, NY 10010-1707.

ISSN: 1078-8956.

DT Article; Journal

FS LIFE; CLIN

LA English

REC Reference Count: 18

*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS.

L6 ANSWER 6 OF 211 USPATFULL

AN 1991:170382 USPATFULL

TI Methods of screening for compounds that derepress or increase telomerase

activity

West, Michael D.; San Carlos, CA, United States

Weinrich, Scott L.; San Francisco, CA, United States

Otani T (Reprint); Nakamura S; Toki M; Motoda R; Kurimoto M; Orita K

AU

PA Dillon, Patrick J., Gaithersburg, MD, United States (U.S.)
 Human Genome Sciences, Inc., Rockville, MD, United States (U.S.)
 corporation) 19990914
 PT US 5932197 19990914
 AI 1997-812003 19970305 (8)
 PRAT US 1996-13609P 19960305 (60)
 DT Utility Granted
 FS
 INL CNT 2123
 INCL INCIM: 435/069, 500
 INCIM: 435/069, 100; 435/252, 300; 435/320, 100; 536/023, 500; 536/024, 300
 NCL NCIM: 435/069, 500
 NCIM: 435/069, 100; 435/252, 300; 435/320, 100; 536/023, 500; 536/024, 300
 IC [6] ICM: C12N015-19
 ICS: C12N015-00; C12N015-63
 EXF 536/23, 5; 435/69, 5; 435/252, 3; 435/320, 1
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 L6 ANSWER 16 OF 211 USPATFULL
 AN 1999-102965 USPATFULL
 TI Mouse model of psoriasis
 Parker, Christiana M., Newton Centre, MA, United States
 Schon, Michael P., Boston, MA, United States
 PA Brigham & Women's Hospital, Inc., Boston, MA, United States (U.S.)
 corporation)
 PI US 545576 19990821
 AI US 1996-628761 19960405 (8)
 DT Utility Granted
 FS
 INL CNT 1935
 INCL INCIM: 800/009, 000
 INCIM: 435/375, 000; 435/377, 000; 424/093, 700; 424/009, 210
 NCL NCIM: 800/009, 000
 NCIM: 424/009, 200; 424/093, 700; 435/375, 000; 435/377, 000
 IC [6] ICM: C12N005-00
 ICS: C12N015-00; A01N063-00; A61K049-00
 EXF 435/375; 435/377; 800/2; 800/DIG.5; 800/DIG.4; 800/9; 424/93, 7; 424/9, 21
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 L6 ANSWER 17 OF 211 USPATFULL
 AN 1999-09279 USPATFULL
 TI Macrophage derived chemokine and chemokine analogs
 IN Goeliski, Ronald, Bothell, WA, United States
 Gray, Patrick W., Seattle, WA, United States
 PA ICOS Corporation, Bothell, WA, United States (U.S. corporation)
 PI US 5932703 19990803
 AI US 1996-66042 19960607 (8)
 RLI Continuation-in-Part of Ser. No. US 1995-558658 filed on 16 Nov 1995
 which is a continuation-in-part of Ser. No. US 1995-479620, filed on 7 Jun 1995
 DT Utility Granted
 FS
 INL CNT 2745
 INCL INCIM: 530/351, 000
 INCIM: 530/324, 000; 930/140, 000; 424/085, 100
 NCL NCIM: 530/351, 000
 NCIM: 424/005, 100; 530/324, 000; 930/140, 000
 IC [6] ICM: C07K014-52
 ICS: A61K038-19
 EXF 530/351; 530/324; 930/140; 424/085, 1
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 L6 ANSWER 18 OF 211 USPATFULL
 AN 1999-085601 USPATFULL
 TI IL-8 receptor antagonists
 IL-8 receptor antagonists
 IN Widdowson, Katherine L., King Prussia, PA, United States
 Nie, Hong, Conshohocken, PA, United States
 Rutledge, Jr., Melvin Clarence, Thousand Oaks, CA, United States
 SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S.)
 corporation) 19990727
 PT US 5922250
 AI US 1998-121264 19980723 (9)
 RLI Continuation-in-Part of Ser. No. WO 1998-US1292, filed on 23 Jan 1998
 PRAI US 1997-42830P 19970408 (60)
 US 1997-35990P 19970123 (60)
 DT Utility Granted
 FS
 INL CNT 1462
 INCL INCIM: 548/361, 100
 INCIM: 548/403, 000
 NCL NCIM: 548/361, 100
 IC [6] ICM: A61K031-115
 ICS: C07D21-56
 EXF 548/361, 11; 51/403
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 L6 ANSWER 19 OF 211 USPATFULL
 AN 1999-085471 USPATFULL
 TI Guainilino, Formanidino, amino and related compounds for inhibiting
 osteoclast-mediated bone resorption
 IN Hartman, George D., Lansdale, PA, United States
 Duigan, Mark E., Schwenksville, PA, United States
 Hoffman, William F., Lansdale, PA, United States
 PA Ihle, Nathan C., Mercer Island, WA, United States
 Merck & Co., Inc., Rahway, NJ, United States (U.S. corporation)
 PI US 5920120 19990727
 AI US 1998-11982 19980130 (9)
 RLI Division of Ser. No. US 1998-714097, filed on 26 Sep 1996, now patented,
 Pat. No. US 5741796 which is a continuation-in-part of Ser. No. US
 1994-250218, filed on 27 May 1994, now abandoned
 DT Utility Granted
 FS
 INL CNT 3417
 INCL INCIM: 514/634, 000
 INCIM: 514/567, 000; 514/568, 000; 514/619, 000; 562/430, 000; 562/439, 000;
 NCL NCIM: 564/1084, 000; 564/1170, 000; 564/246, 000; 564/247, 000
 NCIM: 514/634, 000
 NCIM: 514/567, 000; 514/568, 000; 514/619, 000; 562/430, 000; 562/439, 000;
 IC [6] ICM: A61K031-155
 ICS: A61K031-19; C07C257-10; C07C307-02
 EXF 564/1084, 000; 564/1170, 000; 564/246, 000; 564/247, 000
 564/256; 514/300; 514/311; 514/567; 514/568; 514/619; 514/634; 544/332;
 544/333; 546/164; 546/176; 546/177; 546/668, 1; 546/290; 546/304;
 562/430; 562/439; 564/84; 564/170; 564/246; 564/247
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 L6 ANSWER 20 OF 211 USPATFULL
 AN 1999-05188 USPATFULL
 TI Polynucleotides encoding chemokine alpha-2
 NI Ni, Jian, Rockville, MD, United States
 Gentz, Reiner L., Silver Spring, MD, United States
 Su, Jeffrey Y., Gaithersburg, MD, United States
 Li, Haodong, Gaithersburg, MD, United States
 PA Human Genome Sciences, Inc., Rockville, MD, United States (U.S.)
 corporation)

PI US 5910431 19990608
AI US 1997-825556 19970319 (8)
DT Utility Granted
FS
LN.CNT 2491
INCL INCLM: 435/069.500
INCLS: 435/252.300; 435/320.100; 536/023.500; 536/024.300; 536/024.330
NCL NCLM: 435/069.500
NCLS: 435/252.300; 435/320.100; 536/023.500; 536/024.300; 536/024.330
IC ICM: C12N001-00
ICS: C12N015-00
EXF 536/23.5; 536/4.3-24.33; 435/69.5; 435/752.3; 435/320.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L1: Entry 1 of 1

File: USPT

Nov 9, 1999

US-PAT-NO: 5980887
DOCUMENT-IDENTIFIER: US 5980887 A

TITLE: Methods for enhancing angiogenesis with endothelial progenitor cells

DATE-ISSUED: November 9, 1999

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Isner; Jeffrey M.	Weston	MA		
Asahara; Takayuki	Arlington	MA		

US-CL-CURRENT: 424/93.7; 424/85.1, 424/85.2, 514/44, 514/8

CLAIMS:

What is claimed is:

1. A method for inducing the formation of new blood vessels in an ischemic tissue in a patient in need thereof, comprising:

administering to said patient host an effective amount of an isolated endothelial progenitor cell to induce new blood vessel formation in said ischemic tissue, wherein said endothelial progenitor cell are CD34.sup.+ , flk-1.sup.+ or tie-2.sup.+ .

2. The method of claim 1, further comprising the step of administering to the patient an endothelial cell mitogen or a nucleic acid encoding an endothelial cell mitogen.

3. The method of claim 2, wherein the endothelial cell mitogen is selected from the group consisting of acidic and basic fibroblast growth factors, vascular endothelial growth factor, epidermal growth factor, transforming growth factor α and β ., platelet-derived endothelial growth factor, platelet-derived growth factor, tumor necrosis factor α ., hepatocyte growth factor, insulin like growth factor, erythropoietin, colony stimulating factor, macrophage-CSF, granulocyte/macrophage CSF and nitric oxidesynthase.

4. The method of claim 3, wherein the endothelial cell mitogen is vascular endothelial growth factor.

5. The method of claim 1, wherein said patient is in need of treatment for cerebrovascular ischemia, renal ischemia, pulmonary ischemia, limb ischemia, ischemic cardiomyopathy and myocardial ischemia.

6. A method of enhancing blood vessel formation in a patient in need thereof, comprising:

a. selecting the patient in need thereof;

b. isolating endothelial progenitor cells from the patient, wherein said endothelial progenitor cell are CD34.sup.+ , flk-1.sup.+ or tie-2.sup.+ ; and

c. readministering the endothelial progenitor cells to the patient.

7. A method for treating an injured blood vessel in a patient in need thereof, comprising:

a. selecting the patient in need thereof; and

b. isolating endothelial progenitor cells from the patient, wherein said endothelial progenitor cell are CD34.sup.+ , flk-1.sup.+ or tie-2.sup.+ ; and

c. readministering the endothelial progenitor cells to the patient.

8. The method of claim 7, wherein the injury is the result of balloon angioplasty.

9. The method of claim 7, wherein the injury is the result of deployment of an endovascular stent.

10. The method of claim 7, further comprising the step of administering to the patient an endothelial cell mitogen or a nucleic acid encoding an endothelial cell mitogen.

11. The method of claim 10, wherein the endothelial cell mitogen is selected from the group consisting of acidic and basic fibroblast growth factors, vascular endothelial growth factor, epidermal growth factor, transforming growth factor α and β , platelet-derived endothelial growth factor, platelet-derived growth factor, tumor necrosis factor α , hepatocyte growth factor, insulin like growth factor, erythropoietin, colony stimulating factor, macrophage-CSF, granulocyte/macrophage CSF and nitric oxidesynthase.